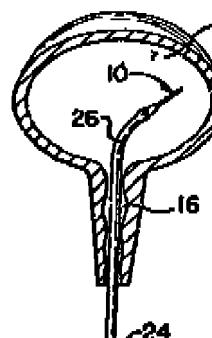
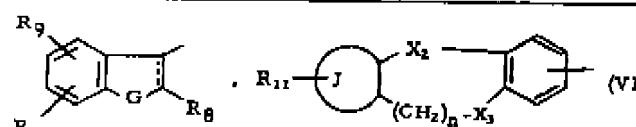
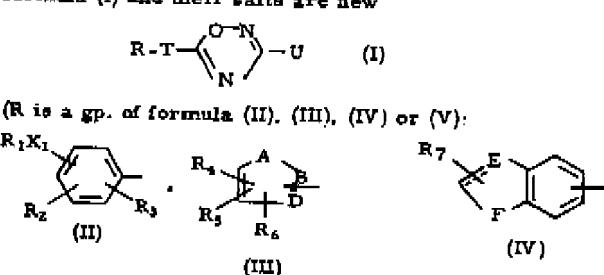


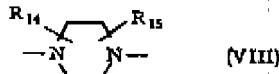
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83-790688/42	B04 D16 J04 K08	INSP 12.03.82 FR 2523-31-A	B(4-B2C, 4-L-D, 4-B4A, 4-B4C, 4-B4D, 4-B4F, 5-A4, 11-C7A, 11-C7B, 12-K4) D(5-Al, 5-H) J(4-B1) 001
INST PASTEUR (CNRS)	12.03.82-FR-004247 (16.09.83) G01n 33/60 C07g 07/00	Aq.- soluble albumin-ligand coupling product - for use in immunoassays	Coupling with albumin increases sensitivity, esp. in the case of enzyme immunoassays for antigens, haptens or antibodies.
C83-102376	Issued in Week 8343. Full Patentee: Inst. Pasteur; Cent. Nat. Rech. Scientifique.	(A) an albumin/specific ligand coupling prod. which is soluble in aq. media is new. (B) Immunoassay of a biological substance (I) comprises: (a) immobilising a substance (II) having binding affinity for (I), (b) incubating with a medium contg. (I), (c) washing the resulting reaction mixt. and incubating with an albumin/specific ligand coupling prod. in soln. in an aq. medium, where the ligand is capable of reacting specifically with (I) or (II), (d) washing the resulting reaction mixt. and incubating with a labelled anti-albumin antibody, and (e) detecting the label. (C) An immunoassay test kit comprises an albumin/specific ligand coupling prod., a labelled anti-albumin antibody and reagents for detecting the label.	DETAILS The specific ligand may be an antigen, hapten, antibody, hormone, hormone receptor, enzyme inhibitor or lectin. It may be coupled with human or animal albumin (esp. BSA) using glutaraldehyde or by 2-stage benzoquinone activation and coupling. The label may be an enzyme, a radioactive material, a fluorochrome, a particulate material or erythrocytes.
	ADVANTAGES		EXAMPLE An BSA/anti-IgE reagent was prep'd. by isolating sheep anti-rabbit Ig antibodies by affinity chromatography, dialysing the antibodies and BSA against phosphate buffer (0.1 M, pH 6.8) at 4°C overnight, and mixing 3 mg of the dialysed antibody with 6 mg of the dialysed BSA in 0.1M phosphate buffer. The mixt. (1 ml) was treated with 0.2 ml of 1% aq. glutaraldehyde and incubated at room temp. for 3 hr. The prod. was used in a sandwich-type enzyme immunoassay for human IgE. (18pp367EDDwgNo0/0). FR 2523311-A
83-795400/43	B07 P34	HEYMANN 18.05.82-US-379480 (+US-353432) (08.09.83) A61m-29	B(11-C4B) 002
	Urological instrument esp. retentive balloon catheter - inserted by sliding over filiform		Pref. the leading section of the filiform (10) is curved as shown. The filiform may be inserted while a stylet wire extends axially within the filiform to stiffen it. Similarly, a stylet tube (24) is placed inside the drainage catheter while it is being slid along the pre-positioned filiform. (25pp295GHDwgNo5/6)
C83-102379	A urological instrument (esp. a catheter) is inserted into the bladder by first advancing a filiform through the urethra, the filiform having smoothly contoured leading end with a lateral opening. Urine flows through this opening and into the filiform to indicate when the leading end of the filiform has entered the bladder. The urological instrument has an internal dia. greater than the external dia. of the filiform to permit the instrument to be slid along the filiform. The instrument may have an inflatable balloon collar which retains the instrument in the bladder; the filiform can then be withdrawn.		
	ADVANTAGE	The correct positioning of the filiform is indicated by the drainage of urine.	AU8311382-A
	EMBODIMENT	Bladder (18) has the drainage catheter (26) in position.	
83-795403/43	B03 (B02)	SUMO 03.03.82 AU8311483-A	B(6-H, 7-E4, 12-D1, 12-D7, 12-D8) 003
SUMITOMO CHEMICAL KK	03.03.82-JP-034168 (08.09.83) A61k-31/41 C07d-271/06 C07d-413/04 C07d-417/10 C07d-471/04 C07d-491/05	5-Aryl-1,2,4-oxadiazole derivs. - are antiinflammatories, analgesics and antipyretics.	
C83-102382	5-Aryl-1,2,4-oxadiazole derivs. of the formula (I) and their salts are new	R ₁ is alkyl, alkenyl, cycloalkyl, cycloalkenyl, opt. substd. phenyl or heterocycl; R ₂ and R ₃ are each H, halo, amino, OH, alkoxy or alkyl; X ₁ is -CH ₂ -; -CH ₂ O-; -CO-; -O-; -S-; -NH or a single bond; R ₄ and R ₅ are H alkyl or opt. substd. phenyl; R ₆ is opt. substd. phenyl or opt. substd. benzoyl; A is N, O or S; B and D are each C or N; E is alkyl, lower alkoxy or opt. substd. phenyl; F is N or C; G is O, S or C or C=C or C=N; broken lines indicate opt. bonds; R ₈ is H or lower alkyl;	
			AU8311483-A

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R_9 is H, halo or alkoxyl;
 R_{10} is H, cyclohexyl or subst. benzoyl;
 G is methylene, subst. benzoylimino, cinnamoylimino or subst. styrylidene, provided that G is $-CH_2-$ when R_{10} is cyclohexyl or subst. benzoyl;
 R_{11} is H, halogen, alkyl or alkoxyl;
 X_2 and X_3 are different and are $-CH_2-$, $-CO-$, $-O-$, $-S-$, $-N^+ - N(CH_3)^+$ or single bond;
 J is a benzene, pyridine, thiophene, furan or pyrrole ring;
 n is 0 or 1;
 T is alkyne or alkenylene each opt. carrying an oxo, OH or lower alkoxyl substit., or T is a single bond;
 U is H, alkyl, alkenyl, polyhaloalkyl, cycloalkyl, cycloalkenyl, opt. subst. phenyl, pyridyl, $-T_1 - R_12$ or $R_{13} - X_4 - T_4 -$; R_{12} is halogen, OH, SH, alkylsulphonyl, dialkoxyimethyl, alkoxycarbonyl, $C(=O)CH_2$, sulpho, CN, $NR' R''$ or $-SR_1 R_1'' X_2 O$;
 R' and R'' are H, alkyl or hydroxy-alkyl;
 $NR' R''$ forms a 5 or 6 membered opt. unsatd. heterocyclic ring, which may contain an O or another N atom, or forms a quaternary ammonium salt or N-oxide;
 R_1' or R_2' are alkyl or alkenyl;
 X is negative monovalent ion;
 T_1 is alkylene or alkenylene, opt. bearing an OXO or OH substit.;

R_{13} is alkyl, alkenyl, hydroxalkyl, acyloxalkyl, aminoalkyl, acylaminooalkyl, cycloalkyl, cycloalkenyl, opt. subst. phenyl, phenyl-alkyl, heterocycl, heterocycl-alkyl, acyl, acylthioalkanoyl, mercaptoalkanoyl, alkoxy-carbonyl, alkylsulphonyl, $-C(=O)R_2 R_2''$ or $SO_2 NR_2 R_2''$;
 R_2' and R_2'' are each H, alkyl or hydroxalkyl;
 X_4 is $-O-$, $-S-$, $-NH-$, a single bond or a gp. of formula (VIII)



R_{14} and R_{15} are each H or alkyl.
All alkyl, alkenyl, alkenylene, cycloalkyl and cycloalkenyl gps. are 'lower' i.e. $\leq 6C$; and cycloalkyl gps. may be oxo- or hydroxy-subst.

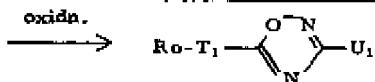
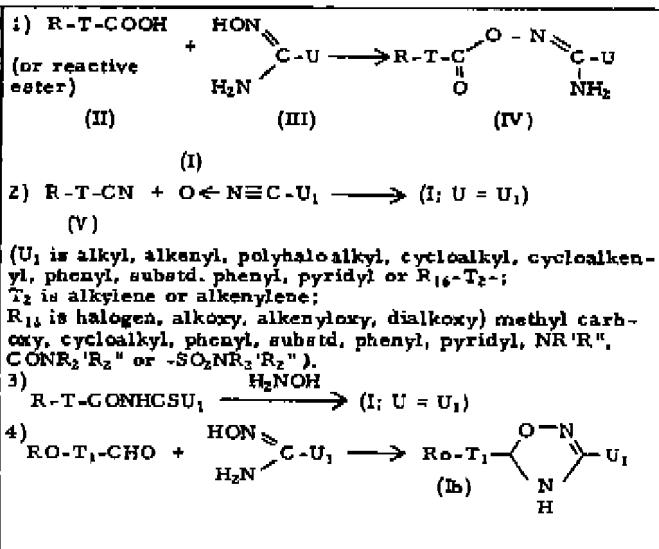
USE

(I) are antiinflammatories, analgesics and antipyretics without ulcerogenic side effects.

PREPARATION

By several methods including:-

AU8311483-A



(R₀ is same as R provided X₁, X₂ and X₃ are not -S-)

EXAMPLE

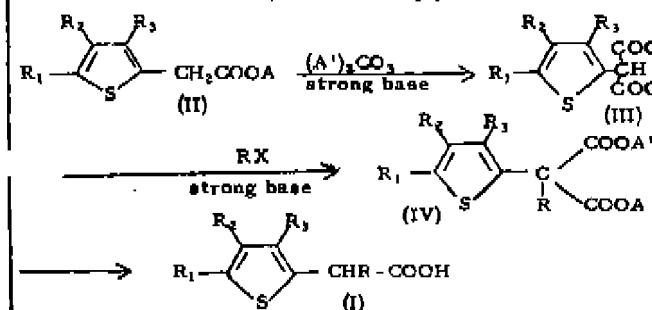
A mixt. of 2-(2-fluoro-4-biphenyl)propionic acid (2.44 g), dry benzene (50 ml) and thionyl chloride (2.38 g) was refluxed for 2 hr., cooled, under reduced pressure and residue dissolved in dry benzene (5 ml). The soln. was added dropwise with cooling to a soln. of acetamidoxime (0.815 g) in dry pyridine and stirred at room temp. and refluxed for 5 hr. The solvent was evapd, under reduced pressure and the residue partitioned between benzene (100 ml) and 10% Na_2CO_3 soln. (20 ml). The organic phase was washed, dried and evapd. and the residue chromatographed on silica gel and eluted with benzene to give 5-(3-fluoro-4-phenyl- α -methylbenzyl)-3-methyl-1,2,4-oxadiazole which was recrystallised from n-hexane to give product (m.p. 55-56°C). (99pp916EDDwgNo0/0).

83-795403/43(3)

AU8311483-A

83-795432/43 803 ROUS 03.12.82
ROUSSEL UCLAF *BE 896-439-A
03.12.82-FR-020271 (12.10.83) C07d
Alpha-alkyl 2-thienyl acetic acid derivs. prodn. - by reacting 2-thienyl acetic acid with alkyl carbonate alkylating agent, then decarboxylation

CB3-102391 (1) Prodn. of α -alkyl- 2-thienylacetic acid derivs. of formula (I) by the following process is new:



B(7-81) + 0 0 4

(R is 1-4C alkyl);
 R_1 , R_2 and R_3 are each H, 1-4C alkyl or halo;
A and A' are 1-4C alkyl; and
X is a functional gp.

(2) The 2-(1,1-di(alkoxycarbonyl)-alkyl)-thiophene intermediates of formula (IV) are new cpds.

USE

(I) are intermediates for pharmaceuticals, esp. anti-inflammatories.

ADVANTAGES

The process uses fewer stages than known methods.

DETAILS

The first stage is pref. in presence of Na ethoxide (esp. 1-1.5 equiv. per mole (II)) at 90-135°C. Reaction of (IV) is esp. also in presence of Na ethoxide, at 50-80°C.

The final stage is by hydrolysis with base, esp. at 50°C to reflux, then acidification with HCl.

The method is esp. used to make (I) where $R_1 = R_2 = R_3 = H$ and R = methyl, esp. (Ia).

EXAMPLE

BE,896439 -A